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Amendments to the Claims:

This listing of claims replaces all prior versions and listings of claims in the application:

Listing of Claims:

- 1-5. (Canceled)
- 6. (Withdrawn) An antibody obtained by the method of claim 1.
- 7. (Withdrawn) A method for producing an antibody with agonistic activity, which comprises the steps of:
- (a) determining the binding activity of an antibody and selecting an antibody with binding activity;
 - (b) modifying the antibody selected in step (a);
- (c) determining the agonistic activity of the modified antibody of step (b) and selecting an antibody with agonistic activity;
- (d) introducing a host cell with a vector carrying a DNA that encodes the antibody selected in step (c); and
 - (e) culturing the host cell of step (d).
- 8. (Withdrawn) The production method of claim 7, wherein the modified antibody is a minibody.
 - 9. (Withdrawn) The production method of claim 8, wherein the minibody is an sc(Fv)2.
- 10. (Withdrawn) The production method of claim 7, wherein the agonistic activity is not determined prior to antibody modification.

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11. (Withdrawn) The production method of claim 7, wherein the antibody is one against a protein expressed on a cell membrane.

12-14. (Canceled)

- 15. (Currently amended) A method of screening comprising:
- (a) providing a plurality of <u>different</u> whole antibodies that bind to a given antigen, wherein the plurality of <u>different</u> whole antibodies comprises antibodies with weak or undetectable agonistic activity for the antigen;
 - (b) producing a minibody form of each whole antibody;
 - (c) screening the minibodies for their ability to agonize the antigen; and
- (d) selecting a minibody if it exhibits agonistic activity greater than that of its respective whole antibody.
- 16. (Previously presented) The method of claim 15, wherein the antigen is a protein expressed on a cell membrane.
 - 17. (Previously presented) The method of claim 15, wherein the antigen is a receptor.
- 18. (Previously presented) The method of claim 15, wherein the antigen is selected from the group consisting of erythropoietin (EPO) receptors, granulocyte colony-stimulating factor (G-CSF) receptors, insulin receptors, Flt-3 ligand receptors, platelet-derived growth factor (PDGF) receptors, interferon (IFN)-α and –β receptors, leptin receptors, growth hormone (GH) receptors, interleukin (IL)-10 receptors, insulin-like growth factor (IGF)-I receptors, leukemia inhibitory factor (LIF) receptors, ciliary neurotrophic factor (CNTF) receptors, HLA-A, HLA-B, HLA-C, HLA-E, HLA-F, HLA-G, HLA-H, HLA-DR, HLA-DQ, HLA-DP, CD1, CD2, CD3, CD4, CD5, CD6, CD7, CD8, CD10, CD11a, CD11b, CD11c, CD13, CD14, CD15s, CD16, CD18, CD19, CD20, CD21, CD23, CD25, CD28, CD29, CD30, CD32, CD33, CD34, CD35, CD38, CD40, CD41a, CD41b, CD42a, CD42b, CD43, CD44, CD45, CD45RO, CD48, CD49a, CD49b, CD49c, CD49d, CD49e, CD49f, CD51, CD54, CD55, CD56, CD57, CD58, CD61,

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CD62E, CD62L, CD62P, CD64, CD69, CD71, CD73, CD95, CD102, CD106, CD122, CD126, and CDw130.

- 19. (Previously presented) The method of claim 15, wherein the antigen is a thrombopoietin (TPO) receptor.
 - 20. (Previously presented) The method of claim 15, wherein the antigen is CD47.
 - 21. (Previously presented) The method of claim 15, wherein the minibodies are sc(Fv)2.
- 22. (Previously presented) The method of claim 15, wherein the minibodies are diabodies.
- 23. (Currently amended) The method of claim 15, wherein the agonistic activities of the different whole antibodies are not assayed prior to step (b).
- 24. (Currently amended) The method of claim 15, wherein the plurality of <u>different</u> whole antibodies are together in a mixture.